## Amendments to the claims:

This listing of claims replaces all prior versions, and listings, of claims in the application.

## **Listing of claims:**

Claims 1-11 (canceled).

Claim 12 (currently amended): A method for measuring an induced release of <sup>13</sup>CO<sub>2</sub> comprising:

- measuring the release of <sup>13</sup>CO<sub>2</sub> in exhaled air of a subject, followed by
- inducing the release of <sup>13</sup>CO<sub>2</sub> in exhaled air of a <u>the</u> subject by intravenous administration of secretin and oral administration of a <sup>13</sup>C-triglyceride to the subject, and
- measuring the release of <sup>13</sup>CO<sub>2</sub> in the exhaled air of the subject before and after the intravenous administration of secretin and before and after the oral administration of the <sup>13</sup>C-triglyceride to the subject.
- Claim 13 (previously presented): The method according to claim 12 characterized in that the <sup>13</sup>C-triglyceride is the mixed triglyceride glyceryl-1,3-dioctadecanoate-2-octanoate-1-<sup>13</sup>C.

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- Claim 14 (previously presented): The method according to claim 12 characterized in that measuring the amount of <sup>13</sup>CO<sub>2</sub> is effected by isotope ratio mass spectrometry (IRMS) or non-dispersive infrared spectroscopy (NDIR).
- Claim 15 (previously presented): The method according to claim 12 wherein the intravenous administration comprises intravenously administering to the subject 1 clinical unit (U) of secretin per kilogram of body weight of the subject within about 15 to 30 minutes.
- Claim 16 (previously presented): The method according to claim 13 wherein the oral administration comprises orally administering to the subject 200 mg of the mixed triglyceride with a test meal.
- Claim 17 (currently amended): A method for diagnosing exocrine pancreatic insufficiency (EPI), comprising:
  - measuring <sup>13</sup>CO<sub>2</sub> in exhaled air of a subject, followed by
  - inducing the release of <sup>13</sup>CO<sub>2</sub> in exhaled air of a <u>the</u> subject by intravenous administration of secretin and oral administration of a <sup>13</sup>C-triglyceride to the subject,
  - measuring the induced value of <sup>13</sup>CO<sub>2</sub> in the exhaled air of the subject before and after intravenous administration of secretin and before and after oral administration of the <sup>13</sup>C-triglyceride to the subject, and

comparing (i) the measured induced value of <sup>13</sup>CO<sub>2</sub> in exhaled air of the subject with
(ii) a measured induced value of <sup>13</sup>CO<sub>2</sub> in exhaled air of a healthy subject after intravenous administration of secretin and after oral administration of the
<sup>13</sup>C-triglyceride to the healthy subject,

wherein a diagnosis of EPI in the subject is indicated when the an induced value of <sup>13</sup>CO<sub>2</sub> in the subject that is reduced as compared to the healthy subject indicates a diagnosis of EPI in the subject.

- Claim 18 (previously presented): The method according to claim 17 characterized in that the <sup>13</sup>C-triglyceride is the mixed triglyceride glyceryl-1,3-dioctadecanoate-2-octanoate-1-<sup>13</sup>C.
- Claim 19 (previously presented): The method according to claim 17 characterized in that measuring the amount of <sup>13</sup>CO<sub>2</sub> is effected by isotope ratio mass spectroscopy (IRMS) or non-dispersive infrared spectroscopy (NDIR).
- Claim 20 (previously presented): The method according to claim 17 wherein the intravenous administration comprises intravenously administering to the subject 1 clinical unit (U) of secretin per kilogram of body weight of the subject within about 15 to 30 minutes.

Claim 21 (previously presented): The method according to claim 20 wherein the oral administration comprises orally administering to the subject 200 mg of the mixed triglyceride with a test meal.